

RNA Editing Technologies Market - Global Industry Size, Share, Trends, Opportunity, and Forecast, Segmented By Type of RNA (Messenger RNA Modification, Transfer RNA Modification, Ribosomal RNA Modification), By Application (Cancer, Genetic Disorders, Temporary Maladies), By Technique (Insertion/ Deletion, Deamination), By Region and Competition, 2019-2029F

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Abstracts

Global RNA Editing Technologies Market was valued at USD 261.28 Million in 2023 and is anticipated to project impressive growth in the forecast period with a CAGR of 6.94% through 2029. The Global RNA Editing Technologies Market encompasses a diverse array of innovative tools and techniques designed to precisely modify RNA molecules for therapeutic, research, and diagnostic applications. RNA editing technologies enable targeted alterations to RNA sequences, offering unprecedented opportunities to correct genetic mutations, modulate gene expression, and develop novel RNA-based therapies. Key players in the market are advancing technologies such as CRISPR-based RNA editing, RNA interference (RNAi), and RNA base editing to address a wide range of diseases, including genetic disorders, cancer, and infectious diseases. The market is driven by increasing research investments, growing demand for personalized medicine, and the expanding applications of RNA editing technologies across various fields of biotechnology and medicine. With ongoing advancements in technology and increasing understanding of RNA biology, the Global RNA Editing Technologies Market is poised for significant growth and innovation in the coming years.

Key Market Drivers



Advancements in CRISPR-Based RNA Editing

CRISPR (Clustered Regularly Interspaced Short Palindromic Repeats) technology has heralded a paradigm shift in the realm of RNA editing, fundamentally altering our ability to precisely modify RNA sequences. This revolutionary technology empowers researchers and scientists with unprecedented control over RNA molecules, enabling targeted alterations with unrivaled precision. Among the arsenal of CRISPR-based tools, CRISPR-Cas13 and CRISPR-CasRx stand out for their exceptional specificity and efficiency in homing in on RNA targets for editing. The advent of CRISPR-based RNA editing has not only transformed basic research but has also catalyzed breakthroughs in therapeutic development, diagnostic applications, and functional genomics. By harnessing the power of CRISPR technology, researchers can manipulate RNA sequences with unparalleled accuracy, opening new avenues for addressing a myriad of diseases at the genetic level.

As the field of RNA editing continues to evolve, ongoing research endeavors are dedicated to refining and enhancing CRISPR-based RNA editing platforms. Efforts are focused on optimizing delivery mechanisms to ensure efficient and precise targeting of RNA molecules within cells. Researchers are striving to expand the scope of CRISPR-based RNA editing by augmenting its targeting capabilities and improving its versatility. These advancements hold tremendous promise for the future of therapeutic interventions, particularly in the realms of genetic disorders and infectious diseases. The enhanced precision and versatility of CRISPR-based RNA editing technologies are anticipated to unlock novel therapeutic strategies, offering hope for patients afflicted with previously untreatable conditions.

Rapid Expansion of RNA Therapeutics

The field of RNA therapeutics has witnessed significant growth in recent years, fueled by advances in RNA editing technologies. RNA-based therapeutics, including mRNA vaccines, RNA interference (RNAi) therapeutics, and RNA editing therapies, hold immense promise for treating a wide range of diseases, from cancer and genetic disorders to infectious diseases and neurological conditions.

RNA editing technologies play a crucial role in the development of RNA therapeutics by enabling precise modifications to RNA sequences to correct disease-causing mutations or modulate gene expression. As the clinical pipeline of RNA therapeutics continues to expand, driven by the success of mRNA vaccines and the growing understanding of



RNA biology, the demand for RNA editing technologies is expected to surge.

Growing Investments in RNA Editing Research

The increasing interest and investment in RNA editing research from both public and private sectors are driving innovation and technological advancements in the field. Funding agencies, academic institutions, biotechnology companies, and pharmaceutical firms are allocating substantial resources to support RNA editing research initiatives, including basic research, preclinical studies, and clinical trials.

These investments are fueling the development of novel RNA editing technologies, the exploration of new therapeutic applications, and the translation of RNA editing-based therapies from the bench to the bedside. Collaborations and partnerships between academia and industry are fostering knowledge exchange, resource sharing, and technology transfer, accelerating the pace of innovation in RNA editing technologies.

Expansion of Genome Editing Applications

While CRISPR-based genome editing initially focused on DNA manipulation, recent advancements have extended its utility to RNA editing applications. Genome editing technologies, such as base editors and prime editors, can be repurposed for RNA editing by targeting RNA molecules instead of genomic DNA.

The versatility of genome editing platforms in RNA manipulation opens up new avenues for RNA-based therapies, gene regulation studies, and functional genomics research. The ability to edit RNA sequences directly offers advantages in terms of temporal control, reversibility, and reduced off-target effects compared to DNA editing approaches. As a result, the expansion of genome editing applications to include RNA editing is driving the growth of the RNA editing technologies market.

Key Market Challenges

Off-Target Effects and Specificity Concerns

A major challenge in the development and adoption of RNA editing technologies is the potential for off-target effects and concerns regarding specificity. Despite advancements in CRISPR-based RNA editing tools, such as CRISPR-Cas13 and CRISPR-CasRx, off-target editing events can still occur, leading to unintended modifications in RNA sequences. These off-target effects pose safety risks and may result in unforeseen



consequences, including cellular toxicity and genetic instability. Addressing off-target effects requires ongoing research efforts to enhance the specificity of RNA editing platforms and minimize unintended alterations to non-target RNA molecules. Strategies such as engineering more precise guide RNAs, optimizing delivery methods, and implementing bioinformatics tools for off-target prediction and validation are being explored to mitigate this challenge. Improving our understanding of RNA biology and the mechanisms underlying off-target editing events is essential for developing safer and more reliable RNA editing technologies.

Precision and Control in RNA Editing

Achieving precise and controlled RNA editing presents a significant challenge in the global RNA editing technologies market. While CRISPR-based RNA editing tools offer remarkable precision, achieving single-nucleotide resolution and site-specific modifications remains challenging. Controlling the timing and duration of RNA editing activity in vivo is crucial for minimizing off-target effects and maximizing therapeutic efficacy.

Strategies to enhance precision and control in RNA editing include the development of novel CRISPR variants with improved editing fidelity, the optimization of guide RNA design to increase specificity, and the implementation of inducible or regulatable editing systems for temporal control. Leveraging insights from RNA biology and RNA-protein interactions can inform the design of RNA editing tools that exhibit greater selectivity and predictability in targeting RNA molecules.

Key Market Trends

Emergence of RNA Editing in Diagnostics

RNA editing technologies are increasingly being utilized in diagnostic applications for detecting and characterizing RNA modifications associated with disease states. RNA biomarkers, such as RNA editing events, alternative splicing patterns, and expression levels of non-coding RNAs, hold diagnostic value for various diseases, including cancer, neurodegenerative disorders, and viral infections.

Advances in RNA sequencing technologies, bioinformatics tools, and machine learning algorithms have facilitated the identification and analysis of RNA biomarkers in clinical samples. RNA editing signatures detected in patient-derived RNA sequences can provide valuable insights into disease progression, treatment response, and patient



prognosis. Consequently, the integration of RNA editing technologies into diagnostic workflows is driving innovation in precision medicine and personalized healthcare.

Regulatory Support for RNA-Based Therapies

Regulatory agencies worldwide are recognizing the potential of RNA-based therapies, including RNA editing technologies, in addressing unmet medical needs and improving patient outcomes. Efforts to streamline regulatory pathways, provide guidance on product development, and facilitate expedited approvals for RNA therapeutics are fostering a favorable environment for innovation and investment in the RNA editing technologies market.

Regulatory frameworks, such as the U.S. Food and Drug Administration's (FDA) RNAi Guidance and the European Medicines Agency's (EMA) guidelines on RNA-based therapeutics, offer clear pathways for the development and regulatory approval of RNA editing-based therapies. These regulatory initiatives provide industry stakeholders with confidence and certainty in navigating the regulatory landscape, accelerating the translation of RNA editing technologies from research to clinical applications.

Segmental Insights

Type of RNA Insights

Based on the type of RNA, Messenger RNA (mRNA) Modification is currently dominating the RNA Editing Technologies Market. mRNA plays a central role in the process of gene expression by conveying genetic information from the DNA to the ribosomes, where it is translated into proteins. RNA modifications, including methylation, and adenosine deamination, regulate mRNA stability, splicing, localization, and translation efficiency, influencing cellular processes and gene expression dynamics.

The emergence of mRNA modification as a dominant force in the RNA Editing Technologies Market is driven by several factors. mRNA modifications are increasingly recognized as critical regulators of gene expression and cellular function, with implications for various physiological and pathological processes, including development, cancer, and neurodegenerative diseases. As researchers unravel the complexity of mRNA modifications and their impact on cellular biology, there is growing interest in developing technologies to manipulate and study these modifications in a controlled manner. mRNA modification technologies offer promising opportunities for therapeutic intervention in the treatment of genetic disorders, cancer, and other



diseases. RNA editing tools, such as CRISPR-based RNA editors and RNA-modifying enzymes, enable precise modifications to mRNA sequences, allowing for the correction of disease-causing mutations, modulation of gene expression, and targeted delivery of therapeutic payloads. These advancements hold potential for the development of RNAbased therapies that harness the power of mRNA modifications to restore normal cellular function and treat a wide range of diseases at the molecular level.

Application Insights

Based on the application segment, cancer stands out as the dominant focus in the Global RNA Editing Technologies Market. The pervasiveness and complexity of cancer have fuelled extensive research efforts aimed at understanding the molecular mechanisms underlying oncogenesis and developing innovative therapeutic strategies. RNA editing technologies offer promising avenues for precision medicine approaches in cancer treatment by targeting aberrant RNA molecules associated with tumour growth, metastasis, and therapy resistance.

The dominance of cancer in driving the Global RNA Editing Technologies Market is underscored by several key factors. RNA editing alterations are increasingly recognized as critical drivers of oncogenesis and tumor progression. Dysregulated RNA editing enzymes, such as ADARs (adenosine deaminases acting on RNA) and APOBECs (apolipoprotein B mRNA editing enzyme, catalytic polypeptide-like), contribute to the generation of cancer-associated RNA editing signatures, including A-to-I (adenosine-toinosine) and C-to-U (cytidine-to-uridine) editing events, which modulate gene expression, RNA stability, and protein function in cancer cells. RNA editing technologies offer innovative approaches for targeting cancer-specific RNA alterations and vulnerabilities. CRISPR-based RNA editing tools, RNA-modifying enzymes, and RNAtargeting small molecules enable precise manipulation of RNA molecules implicated in cancer pathogenesis, providing opportunities for therapeutic intervention. RNA editing strategies can be tailored to correct cancer-driving mutations, disrupt oncogenic RNA modifications, or modulate the expression of key cancer-related genes, thereby suppressing tumor growth and enhancing sensitivity to conventional cancer therapies.

Regional Insights

North America emerges as the dominant region in the Global RNA Editing Technologies Market. Several factors contribute to North America's dominance in this market segment. North America boasts a robust research and development infrastructure, comprising renowned academic institutions, biotechnology hubs, and pharmaceutical



companies at the forefront of RNA editing technologies. These institutions drive innovation and technological advancements in RNA editing research, fostering a dynamic ecosystem for collaboration and discovery. North America benefits from significant investments in healthcare and biotechnology research, supported by government funding agencies, private foundations, and venture capital firms. These investments fuel the development of novel RNA editing technologies, the translation of research findings into clinical applications, and the commercialization of RNA editingbased therapies and diagnostics.

Regulatory frameworks in North America provide clear pathways for the development, approval, and commercialization of RNA editing technologies. Regulatory agencies such as the U.S. Food and Drug Administration (FDA) and Health Canada have established guidelines and expedited review processes for RNA editing-based therapies, facilitating market access and adoption.

Key Market Players

ProQR Therapeutics NV

Sirna Therapeutics, Inc.

Shape Therapeutics, Inc.

Korro Bio, Inc.

Beam Therapeutics, Inc.

QIAGEN N.V.

UniQure NV (UniQure)

LAVA Therapeutics N.V.

argenx SE

Arrowhead Pharmaceuticals, Inc.

Report Scope:



In this report, the Global RNA Editing Technologies Market has been segmented into the following categories, in addition to the industry trends which have also been detailed below:

RNA Editing Technologies Market, By Type of RNA:

oMessenger RNA Modification

oTransfer RNA Modification

oRibosomal RNA Modification

RNA Editing Technologies Market, By Application:

oCancer

oGenetic Disorders

oTemporary Maladies

RNA Editing Technologies Market, By Technique:

olnsertion/ Deletion

oDeamination

RNA Editing Technologies Market, By Region:

oNorth America

United States

Canada

Mexico

oEurope

RNA Editing Technologies Market - Global Industry Size, Share, Trends, Opportunity, and Forecast, Segmented By...



France

United Kingdom

Italy

Germany

Spain

oAsia-Pacific

China

India

Japan

Australia

South Korea

oSouth America

Brazil

Argentina

Colombia

oMiddle East Africa

South Africa

Saudi Arabia



UAE

Competitive Landscape

Company Profiles: Detailed analysis of the major companies present in the Global RNA Editing Technologies Market.

Available Customizations:

Global RNA Editing Technologies market report with the given market data, Tech Sci Research offers customizations according to a company's specific needs. The following customization options are available for the report:

Company Information

Detailed analysis and profiling of additional market players (up to five).



Contents

1.PRODUCT OVERVIEW

- 1.1.Market Definition
- 1.2.Scope of the Market
- 1.2.1.Markets Covered
- 1.2.2.Years Considered for Study
- 1.2.3.Key Market Segmentations

2.RESEARCH METHODOLOGY

- 2.1.Objective of the Study
- 2.2.Baseline Methodology
- 2.3.Key Industry Partners
- 2.4. Major Association and Secondary Sources
- 2.5. Forecasting Methodology
- 2.6.Data Triangulation Validations
- 2.7.Assumptions and Limitations

3.EXECUTIVE SUMMARY

- 3.1. Overview of the Market
- 3.2. Overview of Key Market Segmentations
- 3.3. Overview of Key Market Players
- 3.4. Overview of Key Regions/Countries
- 3.5. Overview of Market Drivers, Challenges, Trends

4.VOICE OF CUSTOMER

5.GLOBAL RNA EDITING TECHNOLOGIES MARKET OUTLOOK

- 5.1.Market Size Forecast
 - 5.1.1.By Value
- 5.2. Market Share Forecast

5.2.1.By Type of RNA (Messenger RNA Modification, Transfer RNA Modification, Ribosomal RNA Modification)

- 5.2.2.By Application (Cancer, Genetic Disorders, Temporary Maladies)
- 5.2.3.By Technique (Insertion/ Deletion, Deamination)



5.2.4.By Region 5.2.5.By Company (2023) 5.3.Market Map

6.NORTH AMERICA RNA EDITING TECHNOLOGIES MARKET OUTLOOK

- 6.1.Market Size Forecast
 - 6.1.1.By Value
- 6.2. Market Share Forecast
- 6.2.1.ByType of RNA
- 6.2.2.By Application
- 6.2.3.By Technique
- 6.2.4.By Country
- 6.3.North America: Country Analysis
 - 6.3.1.United States RNA Editing Technologies Market Outlook
 - 6.3.1.1.Market Size Forecast
 - 6.3.1.1.1.By Value
 - 6.3.1.2. Market Share Forecast
 - 6.3.1.2.1.By Type of RNA
 - 6.3.1.2.2.By Application
 - 6.3.1.2.3.By Technique
 - 6.3.2.Canada RNA Editing Technologies Market Outlook
 - 6.3.2.1.Market Size Forecast
 - 6.3.2.1.1.By Value
 - 6.3.2.2.Market Share Forecast
 - 6.3.2.2.1.By Type of RNA
 - 6.3.2.2.2.By Application
 - 6.3.2.2.3.By Technique
 - 6.3.3.Mexico RNA Editing Technologies Market Outlook
 - 6.3.3.1.Market Size Forecast
 - 6.3.3.1.1.By Value
 - 6.3.3.2. Market Share Forecast
 - 6.3.3.2.1.By Type of RNA
 - 6.3.3.2.2.By Application
 - 6.3.3.2.3.By Technique

7.EUROPE RNA EDITING TECHNOLOGIES MARKET OUTLOOK

7.1.Market Size Forecast



- 7.1.1.By Value
- 7.2.Market Share Forecast
- 7.2.1.By Type of RNA
- 7.2.2.By Application
- 7.2.3.By Technique
- 7.2.4.By Country
- 7.3. Europe: Country Analysis
 - 7.3.1.Germany RNA Editing Technologies Market Outlook
 - 7.3.1.1.Market Size Forecast
 - 7.3.1.1.1.By Value
 - 7.3.1.2.Market Share Forecast
 - 7.3.1.2.1.By Type of RNA
 - 7.3.1.2.2.By Application
 - 7.3.1.2.3.By Technique
 - 7.3.2. United Kingdom RNA Editing Technologies Market Outlook
 - 7.3.2.1.Market Size Forecast
 - 7.3.2.1.1.By Value
 - 7.3.2.2.Market Share Forecast
 - 7.3.2.2.1.By Type of RNA
 - 7.3.2.2.2.By Application
 - 7.3.2.2.3.By Technique
 - 7.3.3.Italy RNA Editing Technologies Market Outlook
 - 7.3.3.1.Market Size Forecast
 - 7.3.3.1.1.By Value
 - 7.3.3.2.Market Share Forecasty
 - 7.3.3.2.1.By Type of RNA
 - 7.3.3.2.2.By Application
 - 7.3.3.2.3.By Technique
 - 7.3.4. France RNA Editing Technologies Market Outlook
 - 7.3.4.1.Market Size Forecast
 - 7.3.4.1.1.By Value
 - 7.3.4.2. Market Share Forecast
 - 7.3.4.2.1.By Type of RNA
 - 7.3.4.2.2.By Application
 - 7.3.4.2.3.By Technique
 - 7.3.5.Spain RNA Editing Technologies Market Outlook
 - 7.3.5.1.Market Size Forecast
 - 7.3.5.1.1.By Value
 - 7.3.5.2. Market Share Forecast



7.3.5.2.1.By Type of RNA 7.3.5.2.2.By Application 7.3.5.2.3.By Technique

8.ASIA-PACIFIC RNA EDITING TECHNOLOGIES MARKET OUTLOOK

8.1.Market Size Forecast

- 8.1.1.By Value
- 8.2. Market Share Forecast
- 8.2.1.By Type of RNA
- 8.2.2.By Application
- 8.2.3.By Technique
- 8.2.4.By Country
- 8.3.Asia-Pacific: Country Analysis
 - 8.3.1. China RNA Editing Technologies Market Outlook
 - 8.3.1.1.Market Size Forecast
 - 8.3.1.1.1.By Value
 - 8.3.1.2. Market Share Forecast
 - 8.3.1.2.1.By Type of RNA
 - 8.3.1.2.2.By Application
 - 8.3.1.2.3.By Technique
 - 8.3.2.India RNA Editing Technologies Market Outlook
 - 8.3.2.1.Market Size Forecast
 - 8.3.2.1.1.By Value
 - 8.3.2.2.Market Share Forecast
 - 8.3.2.2.1.By Type of RNA
 - 8.3.2.2.2.By Application
 - 8.3.2.2.3.By Technique
 - 8.3.3.Japan RNA Editing Technologies Market Outlook
 - 8.3.3.1.Market Size Forecast
 - 8.3.3.1.1.By Value
 - 8.3.3.2. Market Share Forecast
 - 8.3.3.2.1.By Type of RNA
 - 8.3.3.2.2.By Application
 - 8.3.3.2.3.By Technique
 - 8.3.4. South Korea RNA Editing Technologies Market Outlook
 - 8.3.4.1.Market Size Forecast
 - 8.3.4.1.1.By Value
 - 8.3.4.2. Market Share Forecast



8.3.4.2.1.By Type of RNA
8.3.4.2.2.By Application
8.3.4.2.3.By Technique
8.3.5.Australia RNA Editing Technologies Market Outlook
8.3.5.1.Market Size Forecast
8.3.5.1.1.By Value
8.3.5.2.Market Share Forecast
8.3.5.2.1.By Type of RNA
8.3.5.2.2.By Application
8.3.5.2.3.By Technique

9.SOUTH AMERICA RNA EDITING TECHNOLOGIES MARKET OUTLOOK

9.1.Market Size Forecast 9.1.1.By Value 9.2.Market Share Forecast 9.2.1.By Type of RNA 9.2.2.By Application 9.2.3.By Technique 9.2.4.By Country 9.3. South America: Country Analysis 9.3.1.Brazil RNA Editing Technologies Market Outlook 9.3.1.1.Market Size Forecast 9.3.1.1.1.By Value 9.3.1.2. Market Share Forecast 9.3.1.2.1.By Type of RNA 9.3.1.2.2.By Application 9.3.1.2.3.By Technique 9.3.2. Argentina RNA Editing Technologies Market Outlook 9.3.2.1.Market Size Forecast 9.3.2.1.1.By Value 9.3.2.2.Market Share Forecast 9.3.2.2.1.By Type of RNA 9.3.2.2.2.By Application 9.3.2.2.3.By Technique 9.3.3.Colombia RNA Editing Technologies Market Outlook 9.3.3.1.Market Size Forecast 9.3.3.1.1.By Value 9.3.3.2. Market Share Forecast



9.3.3.2.1.By Type of RNA9.3.3.2.2.By Application9.3.3.2.3.By Technique

10.MIDDLE EAST AND AFRICA RNA EDITING TECHNOLOGIES MARKET OUTLOOK

- 10.1.Market Size Forecast
- 10.1.1.By Value
- 10.2.Market Share Forecast
- 10.2.1.By Type of RNA
- 10.2.2.By Application
- 10.2.3.By Technique
- 10.2.4.By Country
- 10.3.MEA: Country Analysis
- 10.3.1. South Africa RNA Editing Technologies Market Outlook
 - 10.3.1.1.Market Size Forecast
 - 10.3.1.1.1.By Value
 - 10.3.1.2. Market Share Forecast
 - 10.3.1.2.1.By Type of RNA
 - 10.3.1.2.2.By Application
 - 10.3.1.2.3.By Technique
- 10.3.2. Saudi Arabia RNA Editing Technologies Market Outlook
 - 10.3.2.1.Market Size Forecast
 - 10.3.2.1.1.By Value
 - 10.3.2.2.Market Share Forecast
 - 10.3.2.2.1.By Type of RNA
 - 10.3.2.2.2.By Application
 - 10.3.2.2.3.By Technique
- 10.3.3.UAE RNA Editing Technologies Market Outlook
- 10.3.3.1.Market Size Forecast
- 10.3.3.1.1.By Value
- 10.3.3.2.Market Share Forecast
- 10.3.3.2.1.By Type of RNA
- 10.3.3.2.2.By Application
- 10.3.3.2.3.By Technique

11.MARKET DYNAMICS



11.1.Drivers

11.2.Challenges

12.MARKET TRENDS DEVELOPMENTS

- 12.1.Merger Acquisition (If Any)
- 12.2.Product Launches (If Any)
- 12.3.Recent Developments

13.PORTER'S FIVE FORCES ANALYSIS

- 13.1.Competition in the Industry
- 13.2.Potential of New Entrants
- 13.3. Power of Suppliers
- 13.4. Power of Customers
- 13.5.Threat of Substitute Products

14.COMPETITIVE LANDSCAPE

14.1.ProQR Therapeutics NV 14.1.1.Business Overview 14.1.2.Company Snapshot 14.1.3. Products Services 14.1.4. Financials (As Reported) 14.1.5.Recent Developments 14.1.6.Key Personnel Details 14.1.7.SWOT Analysis 14.2.Sirna Therapeutics, Inc. 14.3.Shape Therapeutics, Inc. 14.4.Korro Bio, Inc. 14.5.Beam Therapeutics, Inc. 14.6.QIAGEN N.V. 14.7.UniQure NV (UniQure) 14.8.LAVA Therapeutics N.V. 14.9.argenx SE

14.10. Arrowhead Pharmaceuticals, Inc.

15.STRATEGIC RECOMMENDATIONS



16.ABOUT US DISCLAIMER



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